## **IN THE CLAIMS**

1. (Original) A compound according to formula I

wherein R1 = -H, -CN, -COO+, -COS+, -COOH, -COSH, -COOR1.1, -COSR1.1, N-phthalimidyl,

wherein R1.1 = -H, Cl-10 alkyl, C1-10 aralkyl or aryl,

wherein R2 = -H, C1-C4 alkyl, -OR1.1, -Hal (-F -Cl, -Br, -J), -NR2.1R2.2, -Am, -O-Am, -S-Am,

wherein R3 = -H, C1-C4 alkyl, -OR1.1, -Hal (-F -Cl, -Br, -J), -NR2.1R2.2, -Am, -O-Am, -S-Am,

wherein R2.1 = -H, Cl-10 alkyl, C1-10 aralkyl or aryl,

wherein R2.2 = -H, Cl-10 alkyl, C1-10 aralkyl or aryl,

wherein R2.1 and R2.2 may be identical or different,

wherein n and m may be identical or different and 0 to 10,

wherein o and p may be identical or different and 0 to 3,

wherein o > 0, if n and m = 0,

wherein R2 and R3 may be identical or different for Cn and/or Cm,

wherein R2 may be identical or different for every Cx = 1 ... n,

wherein R3 may be identical or different for every  $Cy = 1 \dots m$ ,

wherein -Am is an amino acid radical,

wherein q and r = 0 or 1 and identical or different,

wherein  $-O_r$ - and/or  $-O_q$ - may also be replaced by  $-S_r$ - or  $-S_q$ -, resp.,

wherein -NR2.1R2.2 may be replaced by a linear or branched -C1-C20 alkyl, aralkyl or aryl,

wherein a group -CN, -(CO)-CN, -(CO)-O-R1 or -(CO)-R1 or -C-O-R1 may be replaced by -SO<sub>2</sub>-NR2.1R2.2,

or a physiologically well tolerated salt of such a compound.

- 2. (Original) A compound according to claim 1, wherein R1 = -CN.
- 3. (Currently Amended) A compound according to claim 1 or 2, wherein at least one of the R2 exists at least singly as comprises -Am, wherein -Am preferably represents an amino acid radical of an essential amino acid, wherein in particular q = 0 and r = 1 or q = 1 and r = 0 or q = 1 and r = 1, m = 1, m
- 4. (Currently Amended) A compound according to claim 1 or 2, wherein n = 0 = p = 0, wherein m = 0 to 4, wherein R2 = R3 = -H, or for at least one R2, R2 = -AMm, wherein R2.1 = R2.2 = -H, wherein q = 0 and r = 1.
- (Currently Amended) A compound according to claim 1 or 2, wherein m = p = 0, wherein 0 = 1, wherein n = 0 to 4, wherein R2 = H, or for at least one R2, R2 = -AMm, wherein R3 = -H or -Hal in the case Cx = 1, wherein R3 = -H for all Cx = n > 1, wherein R3 = -H, wherein R3 = -H wherein R3 = -H for all R3 = -H for all R3 = -H wherein R3 = -H for all R3 = -H wherein R3 = -H for all R3 = -H
- 6. (Currently Amended) A compound according to claim 1 or 2, wherein m = 1 to 4, wherein n = 0 = p = 0, wherein R2 = H, or for at least one R2, R2 = -AMm, wherein R3 = -H or -Hal in the case Cy = 1, wherein R3 = -H for all Cy = m > 1, wherein R3 = -H, wherein R3 = -H and R3 = -H for all R3 =

- 7. (Currently Amended) A compound according to claim 1 or 2, wherein o = p = 1, wherein m = 0, wherein n = 0 to 4, wherein R2 = R3 = -H, or for at least one R2, R2 = -AMm, wherein R2.1 = R2.2 = -H, wherein q = 0 and r = 1.
- 8. (Currently Amended) A compound according to claim 1 or 2, wherein n = p = 0, wherein o = 1, wherein m = 0 to 4, wherein R2 = R3 = -H, or for at least one R2, R2 = -AMm, wherein R2 = R2 = -H, wherein q = 0 and q = 0 and q = 0.
- 9. (Currently Amended) A compound according to claim 1 or 2, wherein m = p = 0, wherein o = 1, wherein n = 1 to 4, wherein R2 = R3 = -H, or for at least one R2,  $R2 = -AM\underline{m}$ , wherein R2.1 = R2.2 = -H, wherein q = 0 and r = 1.

## 10. Cancelled

- 11. (Currently Amended) The use of a compound according to Claim 1 for preparing a pharmaceutical composition—A method for treating one or several diseases of from the group comprising [["]]cancer, chronic inflammations, asthma, arthritis, osteoarthritis, chronic polyarthritis, rheumatic arthritis, inflammatory bowl disease, degenerative joint diseases, rheumatic diseases with cartilage disorders, sepsis, autoimmune diseases, type I diabetes, Hashimoto thyreoiditis, autoimmune thrombocytopenia, multiple sclerosis, myasthenia gravis, chronically inflammatory intestinal diseases, Crohn's disease, uveitis, psoriasis, collagenoses, Goodpasture syndrome, diseases with disturbed leukocyte adhesion, cachexia, diseases by increased TNF-alpha concentration, diabetes, adiposity, bacterial infections, in particular with including those by antibiotic resistant bacteria (antibiotic)[["]] comprising administering a pharmaceutical composition prepared comprising the compound according to Claim 1.
- 12. (Currently Amended) A pharmaceutical composition, wherein a compound according to Claim 1 is mixed with one or several physiologically well tolerated auxiliary substances and/or

carrier substances and galenically prepared for the local, in particular oral, or systemic, in particular IV, administration comprising intravenous administration.

13. (Currently Amended) A method The use of a compound according to Claim 1 for inhibiting in vivo the glycolysis and/or the glutaminolysis, in particular of pyruvate kinase, asparaginase, serine dehydratases, transaminases, glutamate oxalacetate transaminase, glutamate pyruvate transaminase, glutamate dehydrogenase, malate dehydrogenase, desaminases and/or glutaminases, in particular in prokaryotes and/or eukaryotes comprising administering a pharmaceutical composition comprising the compound according to Claim 1.